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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO
09/964,824	09/27/2001	Steve Horrigan	669290-73	2602
7590 01/26/2004		EXAMINER		
Alan J. Grant, Esq. c/o Carella, Byrne, Bain, Gilfillan,			SMITH, CAROLYN L	
Cecchi, Stewart & Olstein			ART UNIT	PAPER NUMBER
6 Becker Farm Road			1631	
Roseland, NJ 07068			DATE MAILED: 01/26/2004	1

Please find below and/or attached an Office communication concerning this application or proceeding.

Office Action Summary		Application No.	Applicant(s) HORRIGAN, STEVE	
		09/964,824		
		Examiner	Art Unit	
		Carolyn L Smith	1631	
	The MAILING DATE of this communication a	ppears on the cover sheet w	ith the correspondence address	
	or Reply	DIVIO CET TO EVDIDE AN	AONTH/S) EDOM	
THE - External after aft	HORTENED STATUTORY PERIOD FOR REF MAILING DATE OF THIS COMMUNICATION ensions of time may be available under the provisions of 37 CFR of SIX (6) MONTHS from the mailing date of this communication. The period for reply specified above is less than thirty (30) days, a recommendation of the period for reply is specified above, the maximum statutory period reply within the set or extended period for reply will, by state reply received by the Office later than three months after the mained patent term adjustment. See 37 CFR 1.704(b).	I. 1.136(a). In no event, however, may a eply within the statutory minimum of thiod will apply and will expire SIX (6) MOI ute, cause the application to become A	reply be timely filed  rty (30) days will be considered timely.  NTHS from the mailing date of this communication.  BANDONED (35 U.S.C. § 133).	
	Responsive to communication(s) filed on 19	November 2003.		
		is action is non-final.		
,	Since this application is in condition for allow closed in accordance with the practice unde	vance except for formal mat r <i>Ex parte Quayle</i> , 1935 C.I	tters, prosecution as to the merits is D. 11, 453 O.G. 213.	
Disposit	tion of Claims			
4)⊠	Claim(s) <u>1-39 and 41-54</u> is/are pending in th	e application.		
,	4a) Of the above claim(s) 22-39 and 41-52 is		leration.	
5)□	Claim(s) is/are allowed.			
6)⊠	Claim(s) 1-21,53 and 54 is/are rejected.			
-	Claim(s) <u>1, 3-4, 7-12, 15-21, 53, and 54</u> is/a			
8)🖂	Claim(s) 1-39 and 41-54 are subject to restr	iction and/or election require	ement.	
Applicat	tion Papers			
	The specification is objected to by the Exami			
10)	The drawing(s) filed on is/are: a)☐ a			
	Applicant may not request that any objection to the			
	Replacement drawing sheet(s) including the corr			
11)	The oath or declaration is objected to by the	Examiner. Note the attache	ed Office Action or form PTO-152.	
•	under 35 U.S.C. §§ 119 and 120			
a	Acknowledgment is made of a claim for fore    All b	ents have been received. ents have been received in a riority documents have been eau (PCT Rule 17.2(a)).	Application No n received in this National Stage	
13) :	See the attached detailed Office action for a li Acknowledgment is made of a claim for dome since a specific reference was included in the 37 CFR 1.78. a)   The translation of the foreign language	estic priority under 35 U.S.C first sentence of the specific	. § 119(e) (to a provisional application cation or in an Application Data Shee	
14) <u> </u>	Acknowledgment is made of a claim for dome reference was included in the first sentence of	estic priority under 35 U.S.C f the specification or in an A	. §§ 120 and/or 121 since a specific pplication Data Sheet. 37 CFR 1.78.	
Attachme	nt(s)			
1)	ice of References Cited (PTO-892) ice of Draftsperson's Patent Drawing Review (PTO-948) propertion Disclosure Statement(s) (PTO-1449) Paper Note	5) Notice of	Summary (PTO-413) Paper No(s) Informal Patent Application (PTO-152)	

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### **DETAILED ACTION**

Applicant's amendments and remarks, filed 11/19/03, are acknowledged. Amended claims 1, 2, 4, 6, 13, 15-19, and 53-54 are acknowledged. On page 10 (first sentence) of Applicant's response, it is incorrectly noted that claims 1-21, 53, and 54 are pending. Actually, claims 1-39 and 41-54 are pending while claims 22-39 and 41-52 are withdrawn from consideration as being drawn to non-elected Groups.

Applicant's arguments, filed 11/19/03, have been fully considered but they are not deemed to be persuasive. Rejections and/or objections not reiterated from the previous office actions are hereby withdrawn. The following rejections and/or objections are either reiterated or newly applied. They constitute the complete set presently being applied to the instant application.

Claims 1-21, 53, and 54 are herein under examination.

# Claim Objections

Claims 1, 3-4, 7-12, 15-21, 53, and 54 are objected to due to their inclusion of subject matter that has been non-elected due to a restriction requirement and therefore withdrawn from consideration. The non-elected subject matter of claims 1, 3-4, 7-12, 15-21, 53, and 54 is as follows: Claims 1, 3-4, 7-12, 15-21, 53, and 54 contain sequences, such as sequences other than

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SEQ ID NO: 137, 164, 174, 180, 191, 199, 344, 381, 390, and 465, which are non-elected subject matter. Removal of non-elected subject matter is requested.

# Claim Rejections – 35 U.S.C. 112, first paragraph

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

## LACK OF WRITTEN DESCRIPTION

The rejection of claims 1-21, 53, and 54 is maintained under 35 U.S.C. 112 as containing subject matter which was not described in the specification in such a way as to reasonably convey to one skilled in the relevant art that the inventor, at the time of the invention was filed, had possession of the claimed invention.

The specification discloses SEQ ID NO: 137, 164, 174, 180, 191, 199, 344, 381, 390, and 465 which correspond to nucleic acid sequences. SEQ ID NO: 137, 164, 174, 180, 191, 199, 344, 381, 390, and 465 and their full complements meet the written description provisions of 35 U.S.C. 112, first paragraph. However, due to the open claim language of "expressing a gene that corresponds to a polynucleotide" (claim 1) and "comprising a nucleotide sequence corresponding to a gene" (claim 54), these claims encompass sequences which do not meet the written description provision of 35 U.S.C. 112, first paragraph. The specification provides insufficient written description to support the genus encompassed by these claims.

Vas-Cath Inc. v. Mahurkar, 19 USPQ2d 1111, makes clear that "applicant must convey with reasonable clarity to those skilled in the art that, as of the filing date sought, he or she was in possession of the invention. The invention is, for purposes of the 'written description' inquiry, Application/Control Number: 09/964,824

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whatever is now claimed." (See page 1117.) The specification does not "clearly allow persons of ordinary skill in the art to recognize that [he or she] invented what is claimed." (See <u>Vas-Cath</u> at page 1116.)

With the exception of SEQ ID NO: 137, 164, 174, 180, 191, 199, 344, 381, 390, and 465, the skilled artisan cannot envision the detailed chemical structure of the encompassed polynucleotides and/or proteins, regardless of the complexity or simplicity of the method of isolation. Adequate written description requires more than a mere statement that it is part of the invention and reference to a potential method for isolating it. The nucleic acid itself is required. See Fiers v. Revel, 25 USPQ2d 1601, 1606 (CAFC 1993) and Amgen Inc. V. Chugai Pharmacentical Co. Ltd., 18 USPQ2d 1016. In Fiddes v. Baird, 30 USPQ2d 1481, 1483, claims directed to mammalian FGF's were found unpatentable due to lack of written description for the broad class. The specification provided only the bovine sequence.

Finally, <u>University of California v. Eli Lilly and Co.</u>, 43 USPQ2d 1398, 1404, 1405 held that:

...To fulfill the written description requirement, a patent specification must describe an invention and do so in sufficient detail that one skilled in the art can clearly conclude that "the inventor invented the claimed invention." *Lockwood v. American Airlines, Inc.*, 107 F.3d 1565, 1572, 41 USPQ2d 1961, 1966 (1997); *In re Gosteli*, 872 F.2d 1008, 1012, 10 USPQ2d 1614, 1618 (Fed. Cir. 1989) (" [T]he description must clearly allow persons of ordinary skill in the art to recognize that [the inventor] invented what is claimed."). Thus, an applicant complies with the written description requirement "by describing the invention, with all its claimed limitations, not that which makes it obvious," and by using "such descriptive means as words, structures, figures, diagrams, formulas, etc., that set forth the claimed invention." *Lockwood*, 107 F.3d at 1572, 41 USPQ2d at 1966.

Therefore, only SEQ ID NO: 137, 164, 174, 180, 191, 199, 344, 381, 390, and 465 and their full length complements, but not the full breadth of the claims meet the written description provision of 35 USC 112, first paragraph. The species specifically disclosed are not representative of the genus because the genus is highly variant. Applicant is reminded that <u>Vas-Cath</u> makes clear that the written description provision of 35 USC 112 is severable from its enablement provision. (See page 1115.)

Applicant states this rejection cannot be solely based on the use of open ended language because "comprising" is standard claim language and that this rejection must be predicated on

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the use of the term "corresponding" in conjunction with "comprising". It is noted that this rejection is made because these phrases, in their broadest and reasonable interpretation, encompass sequences that do not have written support in the specification, claims, and/or drawings as originally filed. Applicant states the term "correspond" is defined as meaning a gene that encodes a RNA at least 90% identical to the claimed polynucleotide. This is found unpersuasive as this sequence could encompass a claimed sequence, plus up to 10% of additional sequence on either end that does not meet the written description provision of 35 U.S.C. 112, first paragraph. Again, it is acknowledged that the Applicant has written support for SEQ ID NO: 137, 164, 174, 180, 191, 199, 344, 381, 390, and 465 and their full length complements, but not for the full breath of the claims. The arguments presented by the Applicant do not specifically address the written description rejection, as reiterated, and is therefore found unpersuasive.

## Claim Rejections – 35 U.S.C. 112, first paragraph

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

Factors to be considered in determining whether a disclosure would require undue experimentation have been summarized in *Ex parte Forman*, 230 USPQ 546 (BPAI 1986) and reiterated by the Court of Appeals in *In re Wands*, 8 USPQ2d 1400 at 1404 (CAFC 1988). The factors to be considered in determining whether undue experimentation is required include: (1) the quantity of experimentation necessary, (2) the amount or direction presented, (3) the presence or absence of working examples, (4) the nature of the invention, (5) the state of the prior art, (6)

the relative skill of those in the art, (7) the predictability or unpredictability of the art, and (8) the breadth of the claims. The Board also stated that although the level of the skill in molecular biology is high, the results of experiments in genetic engineering are unpredictable. While all of these factors are considered, a sufficient amount for a *prima facie* case are discussed below.

#### LACK OF ENABLEMENT

Claims 1-21, 53, and 54 are rejected under 35 U.S.C. § 112, first paragraph, as containing subject matter which was not described in the specification in such a way as to enable one skilled in the art to which it pertains, or with which it is most nearly connected, to make and/or use the claimed invention.

The sequences selected in these methods include sequences from GenBank with accession numbers AB011095, G42509, AA281006, AA490819, N71063, AA243738, AA411711, N73808, R27957, and H05625. AB011095 is from RNA from human male brain; G42509 is from human genomic DNA; AA281006 is from cDNA from human germinal center B cells; AA490819 is from cDNA from human germinal center B cells; N71063 is from cDNA from human fetal lung; AA243738 is from cDNA from pooled human melanocyte, fetal heart, and pregnant uterus; AA411711 is from cDNA from pooled human melanocyte, fetal heart, and pregnant uterus; N73808 is from cDNA from human male multiple sclerosis legions tissue; R27957 is from cDNA from human female placenta; and H05625 is from cDNA from human female infant brain. There are millions of sequences in the world with a small portion actually available in public databases, such as GenBank. A microarray type of invention that involves a multitude of sequences that appear to be randomly selected with no previously known function or association with cancer that merely come from brain, random genomic DNA, B cells,

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melanocytes, fetal organs, female organs, and multiple sclerosis lesions (as is the case with the elected sequences) do not appear to be enabling for screening chemical compounds for antineoplastic activity. The quantity of experimentation required to verify that these sequences represent valid predictors of screening chemical compounds for anti-neoplastic activity appears to be undue. Due to undue experimentation required, the lack of guidance directed to verifying such sequences functioning as valid predictors, the lack of working examples addressing the same, the unpredictability of knowing if these sequences are potentially valid predictors for screening anti-neoplastic activity, and the breath of the claims; this invention is rejected due to the lack of enablement for one skilled in the art to be able to make and use the invention.

#### Conclusion

No claim is allowed.

Papers related to this application may be submitted to Technical Center 1600 by facsimile transmission. Papers should be faxed to Technical Center 1600 via the PTO Fax Center located in Crystal Mall 1. The faxing of such papers must conform with the notices published in the Official Gazette, 1096 OG 30 (November 15, 1988), 1156 OG 61 (November 16, 1993), and 1157 OG 94 (December 28, 1993) (See 37 CFR §1.6(d)). The CM1 Fax Center number is (703) 872-9306.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Carolyn Smith, whose telephone number is (571) 272-0721. The examiner can normally be reached Monday through Thursday from 8 A.M. to 6:30 P.M.

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If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Michael Woodward, can be reached on (571) 272-0722.

Any inquiry of a general nature or relating to the status of this application should be directed to Legal Instruments Examiner Tina Plunkett whose telephone number is (571) 272-0549 or to the Technical Center receptionist whose telephone number is (703) 308-0196.

January 13, 2004

ARDIN H. MARSCHEL PRIMARY EXAMINER